

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:

Gurtej Singh Sandhu et al.

Title:

Filed:

METHOD TO REDUCE FIXED CHARGE IN CVD OZONE DEPOSITED FILMS

Docket No.:

Examiner:

303.573US1

April 22, 1996

Matthew Wipple

Serial No.: 08/636,069

Due Date: September 15, 1999

Group Art Unit: 2813

Assistant Commissioner for Patents

Washington, D.C. 20231

We are transmitting herewith the following attached items (as indicated with an "X"):

X A return postcard.

X An Amendment and Response (6 Pages).

Please consider this a PETITION FOR EXTENSION OF TIME for sufficient number of months to enter these papers and please charge any additional required fees or credit overpayment to Deposit Account No. 19-0743.

CERTIFICATE UNDER 37 CFR 1.8: The undersigned hereby certifies that this Transmittal Letter and the paper, as described above, are being deposited in the United States Postal Service, as first class mail, in an envelope addressed to: Assistant Commissioner for Patents, Washington, D.C. 20231, on this ______ day of _September_, 1999.

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(GENERAL)

SEP 14 1939

S/N 08/636,069

PATENT

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AMENDMENT AND RESPONSE

Assistant Commissioner for Patents Washington, D.C. 20231

16/0 FJONES 9-29-99

Applicant has reviewed the Office Action mailed June 15, 1999. Please amend the application as follows:

IN THE CLAIMS

Please amend claims 31, 42, 51 and 52 as follows:

31. (Once amended)

A method of depositing a silicon dioxide layer on a substrate

surface, comprising:

contacting the substrate surface with a reaction volume of gas comprising a SiO₂

precursor and ozone; [and]

heating the substrate surface to a temperature of about 480°C to 700°C; and

illuminating the reaction volume of gas from a light source without directly exposing the

substrate surface to the light source.

(Once amended)

A method of depositing a doped silicon dioxide layer on a substrate

surface, comprising:

contacting the substrate surface with a reaction volume of gas comprising a SiO₂

precursor, ozone and at/least one dopant source; [and]

heating the substrate surface to a temperature of about 480°C to 700°C; and

illuminating the reaction volume of gas from a light source without directly exposing the

substrate surface to the light source.

SEP 14 1990

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